NRC Form 483
1-76
10 CFR 31

APPROVED BY GAO
B0-R0100

REGISTRATION CERTIFICATE—IN VITRO TESTING WITH BYPRODUCT MATERIAL UNDER GENERAL LICENSE

Section 31.11 of the U.S. Nuclear Regulatory Commission's regulations [illegible] physicians, clinical laboratories, and hospitals [illegible] possess certain small quantities of byproduct material [illegible] in vitro laboratory tests not involving the internal or external administration of the byproduct material [illegible] to human beings or animals. Possession of byproduct material under 10 CFR 31.11 is not [illegible] physician, clinical laboratory, or hospital has filed NRC Form 483 and received from the [illegible] NRC Form 483 [illegible] registration number.

Carlito Majica
3677 S. Lapeer Rd
Lake Orion, MI 48035

3. I hereby apply for a registration number pursuant to § 31.11, 10 CFR 31 for use of byproduct materials for (*Please check one block only*)
   - [✓] a. Myself, a duly licensed physician authorized to dispense drugs in the practice of medicine.
   - [ ] b. The above-named clinical laboratory.
   - [ ] c. The above-named hospital.
4. To be completed by the Nuclear Regulatory Commission.

Registration number: 6539

FOR THE U. S. NUCLEAR REGULATORY COMMISSION

Shirley A. Crutchfield November 29, 1982

INSTRUCTIONS

1. Submit this form in triplicate to
   Office of Nuclear Material Safety and Safeguards
   ATTN: Radioisotopes Licensing Branch
   U.S. Nuclear Regulatory Commission
   Washington, D.C. 20555
2. Please print or type the name and address (including zip code) of the registrant physician, clinical laboratory, or hospital for whom or for which this registration form is filed. Enter one letter or [illegible] the address below the dotted line [illegible] not extend the address beyond the right margin. (At NRC, a registration number will be assigned and a validated copy of NRC Form 483 will be returned.)

5. If place of use is different [illegible]

6. Certification

I hereby certify:

a. All information in this [illegible]

b. The registrant has applied [illegible] tests for which byproduct material will be used under the general license of 10 CFR [illegible]. The tests will be performed only by trained and competent [illegible] in the use of the instruments and in the handling of the byproduct material. [illegible]

c. I understand that Commission regulations require that any change in the information furnished by a registrant on this registration certificate be reported to the Director of Nuclear Material Safety and Safeguards within 30 days from the effective date of such change.

d. I have read and understand the provisions of Section 31.11 of NRC regulations 10 CFR 31 (reprinted on the reverse side of this form); and I understand that the registrant is required to comply with the [illegible] to all byproduct material which he receives, acquires, possesses, uses, or transfers under the general license for which this Registration Certificate is filed with the Nuclear Regulatory Commission.

Date 10-19-82

By Carlito C. Majica M.D.
Signature of person filing form

Printed name and title of position of person filing form

WARNING—18 U.S.C., Section [illegible] of June [illegible] reprint [illegible] to make a willfully false statement or [illegible] jurisdiction [illegible]